

## *Supporting Information*

# **Electrochemical pinacol coupling of aromatic carbonyl compounds in a [BMIM][BF<sub>4</sub>]/H<sub>2</sub>O mixture**

Hannah Kronenwetter, Jakub Husek, Brian Etz, Aaron Jones, and Renuka Manchanayakage\*

Department of Chemistry, Susquehanna University, 514 University Avenue, Selinsgrove, PA 17870

manchanayakage@susqu.edu

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## General Information

All reactions were conducted under nitrogen atmosphere. All reagents were obtained from commercial suppliers and used without further purification. Ionic liquids, [BMIM][BF<sub>4</sub>] and [BMIM][PF<sub>6</sub>] were synthesized according to the literature.<sup>1</sup> NMR experiments were conducted in chloroform-*d*.

## General procedure for the electrochemical pinacol coupling reaction

The carbonyl compound (0.35 mmol) and a mixture of 80% [BMIM][BF<sub>4</sub>]/H<sub>2</sub>O (4.0 mL of [BMIM][BF<sub>4</sub>] and 1.0 mL of H<sub>2</sub>O) were placed in an electrochemical cell fitted with a tin foil anode (1.0 cm<sup>2</sup>), a platinum plate cathode (1.0 cm<sup>2</sup>) and a Ag/AgCl reference electrode. The mixture was stirred and degassed by bubbling nitrogen for 30 minutes. A controlled potential of 2.0 V was applied under nitrogen atmosphere for 5 hours. The resultant solution was extracted with diethyl ether (2 × 5 mL). The ether extract was dried with anhydrous sodium sulfate, filtered and concentrated in vacuo. When necessary, the residue was purified via silica gel flash column chromatography (10% ethyl acetate in hexane eluent) to afford the final product.

## References

1. Dupont, J.; Consorti, C.; Suarez, P.; Souza, R. *Org. Syn. Coll. Vol.* **2004**, 79, 236.

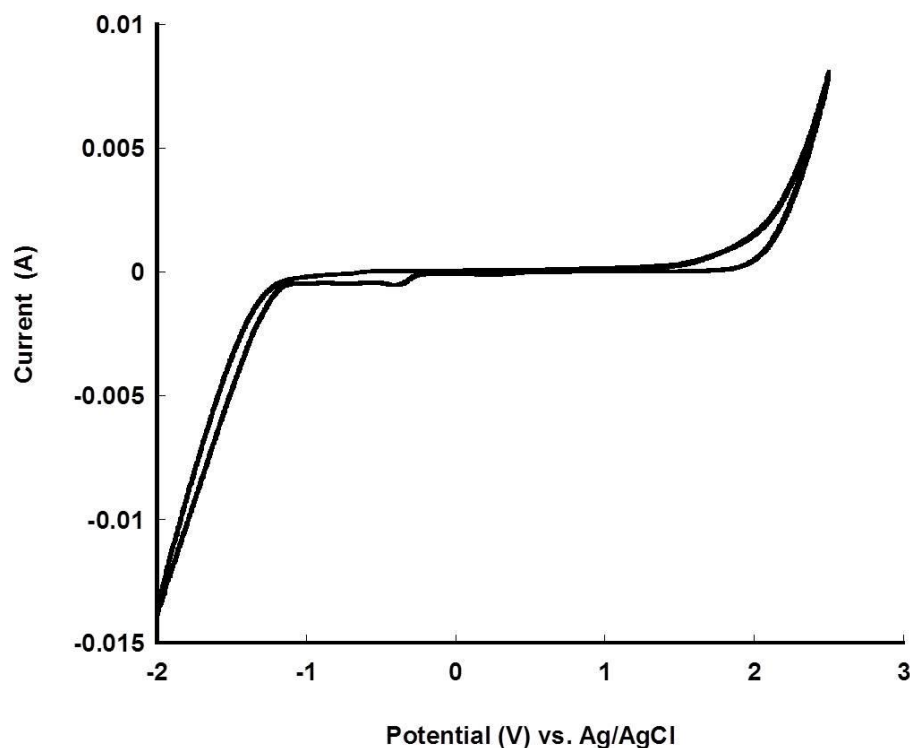


Figure 1: Cyclic voltammogram of the 80% [BMIM][BF<sub>4</sub>]/H<sub>2</sub>O system. Recorded at a Pt cathode against Ag/AgCl reference. The scan rate was 0.025 Vs<sup>-1</sup>.

